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CONFIRMATION NO. ATTORNEY DOCKET NO. FIRST NAMED INVENTOR FILING DATE APPLICATION NO. 8329 A-69566-1/RFT/RMS/RMK Arthur J. Chirino 07/10/2001 09/903,378 04/09/2003 7590 EXAMINER FLEHR HOHBACH TEST ALBRITTON & HERBERT LLP BORIN, MICHAEL L **Suite 3400** Four Embarcadero Center PAPER NUMBER San Francisco, CA 94111-4187 ART UNIT 1631 10 DATE MAILED: 04/09/2003

Please find below and/or attached an Office communication concerning this application or proceeding.



Application No. 09/903,378

Applicant(s)

Chirino et al

Office Action Summary

Examiner

Michael Borin

Art Unit **1631**



	The MAILING DATE of this communication appears of	n the cover sheet with the correspondence address	
Period for Reply A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE			
Status	•		
1) 💢	Responsive to communication(s) filed on Jan 28, 20	003	
2a) □	This action is FINAL . 2b) \(\overline{\text{Z}} \) This acti	ion is non-final.	
3) [Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11; 453 O.G. 213.		
Dispo	sition of Claims	is/are pending in the application.	
4) 💢	Claim(s) <u>1-17</u>	is/are pending in the application.	
	4a) Of the above, claim(s)	is/ale withdrawn from concessor	
5)	Claim(s)	IS/die dilowedi	
6) 🔯	Claim(s) 1-17	is/are rejected.	
-	7 01: (-)	15/4/6 00/00/00 10	
7) L	Claim(s)	are subject to restriction and/or election requirement.	
8) [
Application Papers			
9) L	9) The specification is objected to by the Examiner. 10) The drawing(s) filed on is/are a) accepted or b) objected to by the Examiner.		
The drawing(s) filed on			
12) The oath or declaration is objected to by the Examiner.			
Priority under 35 U.S.C. §§ 119 and 120 13) Acknowledgement is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).			
a) □ All b) □ Some* c) □ None of:			
1 Certified copies of the priority documents have been received.			
Continue application No.			
Copies of the certified copies of the priority documents have been received in this National Stage Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). *See the attached detailed Office action for a list of the certified copies not received.			
14) Acknowledgement is made of a claim for domestic priority under 35 U.S.C. § 119(e).			
The translation of the foreign language provisional application has been received.			
a) Ine translation of the foldight displays provided as Ine translation of the Ine translation of			
1			
	achment(s) Notice of References Cited (PTO-892)	4) Interview Summary (PTO-413) Paper No(s).	
	✓ Notice of Draftsperson's Patent Drawing Review (PTO-948)	5) Notice of Informal Patent Application (PTO-152)	
	☐ Information Disclosure Statement(s) (PTO-1449) Paper No(s)5 ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐	6) Cther:	

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DETAILED ACTION

Status of Claims

1. Response to restriction requirement filed 1/28/03 is acknowledged. Applicant elected, without traverse, Group I, claims 1-17. Claims 19 is canceled. Claims 1-17 are pending.

Claim Rejections - 35 USC § 112, second paragraph.

The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

- 2. Claim 1-17 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.
- A. The claims are drawn to modulating protein immunogenecity which is a biological function determined experimentally. The entire claimed method, however, is *in silico*, i.e. all steps are drawn to computer modeling. It remains vague and indefinite, without an experimental testing of functions of candidate variant protein, whether computer modeling steps result in modulation of immunogenecity.

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B. Claim 1 recites "immunogenecity filter". The term is not defined in the claim. The specification briefly explains that "immunogenecity filter" is "any one of a number of scoring functions" related to binding to different epitopes. What is such scoring function, how it is applied to the candidate protein, and what proteins satisfy the criteria of "modulated immunogenecity" is not clear. Thus, both the nature of the method step, and metes and bounds of the claimed method remain unclear.

C. It is not clear, how the step of optimizing for a scoring function addressed in claims 6-8 is related to step c) of the base claim 1: is it a further step in addition to steps a)-c) of claim 1, a step alternative to step c) of claim 1, or it is the step c) of claim 1.

Claim Rejections - 35 USC § 112, first paragraph.

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

3. Claims 1-17 are rejected under 35 U.S.C. 112, first paragraph, as containing subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventors, at the time the application was filed, had possession of the claimed invention.

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The claims are drawn to modulating of immunogenecity of proteins. There is no single example in the specification of the operability of the method neither *in silico*, nor in experimental conditions on a real protein synthesized following its *in silico* design. The only mention of "immunogenecity filter" on p. 30 (lines 15-21) is so vague that it is not clear whether applicant was in possession of any algorithm or scoring function that would result in a design of a protein with altered immunogenecity.

The inventor must be able to describe the item to be patented with such clarity that the reader is assured that the inventor actually has possession and knowledge of the unique method that makes it worthy of patent protection. The reader can certainly appreciate the goal but establishing goals does not make a patent. As the Court of Appeals for the Federal Circuit stated in a case involving similar issues, an inadequate patent description that merely identifies a plan to accomplish an intended result "is an attempt to preempt the future before it has arrived." Fiers v. Revel, 984 F.2d 1164, 1171 (Fed. Cir.1993). To satisfy the written-description requirement, the specification must describe every element of the claimed invention in sufficient detail so that one of ordinary skill in the art would recognize that the inventor possessed the claimed invention at the time of filing. Vas-Cath, 935 F.3d at 1563; see also Lockwood v. American Airlines, Inc., 107 F.3d 1565, 1572 (Fed. Cir.

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1997) (patent specification must describe an invention and do so in sufficient detail that one skilled in the art can clearly conclude that "the inventor invented the claimed invention"). There is no demonstration in the specification that applicants generated any compound which, after computer generation, and application of "computational immunogenecity filters" had immunogenecity different from that of parent molecule. Similarly to *In re Wilder*, 736 F.2d 1516 (Fed. Cir. 1984), *cert. denied*, 469 U.S. 1209 (1985)the specification did "little more than outline] goals appellants hope the claimed invention achieves and the problems the invention will hopefully ameliorate."

Claim Rejections - 35 USC § 102 and 103.

The following is a quotation of the appropriate paragraphs of 35 U.S.C.102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless --

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

4. Claims 1-17 are rejected under 35 U.S.C. 103(a) as obvious over Fleckenstein et al (Eur. J. Biochem., 240, 71-77, 1996) or Abrams (Current Opinions in Immunology, 12, 85-91, 2000; references C15 and C1, respectively) in view of Altuvia et al or Meister et al or Buus et al (references C2, C37, and C9, respectively) and further in view of Mayo et al (WO 98/47089 or US Patent 6,269,312; references B1 and A1, respectively).

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The instant claims are drawn to method of modulating immunogenecity of a protein comprising the steps of inputting the protein's structure into a computer, modulating the structure at variable positions, and identifying candidate variant proteins by applying "computational immunogenecity filter". The latter "filter", as explained in specification, p. 30, can be any of scoring functions derived on binding of peptides to MHC molecules, or T cell epitopes or B cell epitopes.

Fleckenstein et al (Eur. J. Biochem., 240, 71-77, 1996) teaches method for determining peptides with modulated immunogenecity (i.e., with altered binding to leucocyte antigens to MHC molecules). Peptide libraries of undecapeptides with substitutions at variable positions are prepared synthetically, and binding of the peptides to human leukocyte antigen DRB1 is used as a "immunogenecity filter" to determine variant peptide immunogenecity. Abrams teaches that to modify MHC binding reactivity of peptides, rational targeted substitution of amino acid residues can be introduced to peptide ligands for regulation of immunogenic responses (p. 89). The referenced methods differs from the claimed invention in that both generation of variants and their testing are done in experimental conditions, not *in silico*.

There are numerous publications describing use of computerized algorithms to predict binding of peptides to MHC molecules. See, for example references of Altuvia et al or Meister et al or Buus et al, cited by applicants. Thus, it would have been

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prima facie obvious to one skilled in the art at the time the invention was made to be motivated to substitute experimental determination of the immunogeneoity of the candidate variant peptides with computerized estimates of their immunogeneoity, such as described in Altuvia et al or Meister et al or Buus et al.

Further, in regard to method of generating of candidate peptides, computerized way of generating peptide in the claimed method does not render the referenced methods utilizing chemical preparation of the peptides. Alternatively, computerized methods of generating peptide libraries with substitutions at variable positions proved to be an efficient way of modeling peptides which are further assessed for their biological functions. See for example, Mayo et al (WO 98/47089) or Mayo et al (US Patent 6,269,312).

Conclusion.

- 5. No claims are allowed
- 6. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Michael Borin whose telephone number is (703) 305-4506. Dr. Borin can normally be reached between the hours of 8:30 A.M. to 5:00 P.M. EST Monday to Friday. If attempts to reach the examiner by telephone are

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unsuccessful, the examiner's supervisor, Mr. Michael Woodward, can be reached on (703) 308-4028. The fax telephone number for this group is (703) 305-3014.

Any inquiry of a general nature or relating the status of this application should be directed to the Group receptionist whose telephone number is (703) 308-0196.

March 31, 2003

MICHAEL BORIN, PH.D. PRIMARY EXAMINER

mlb

Mpm